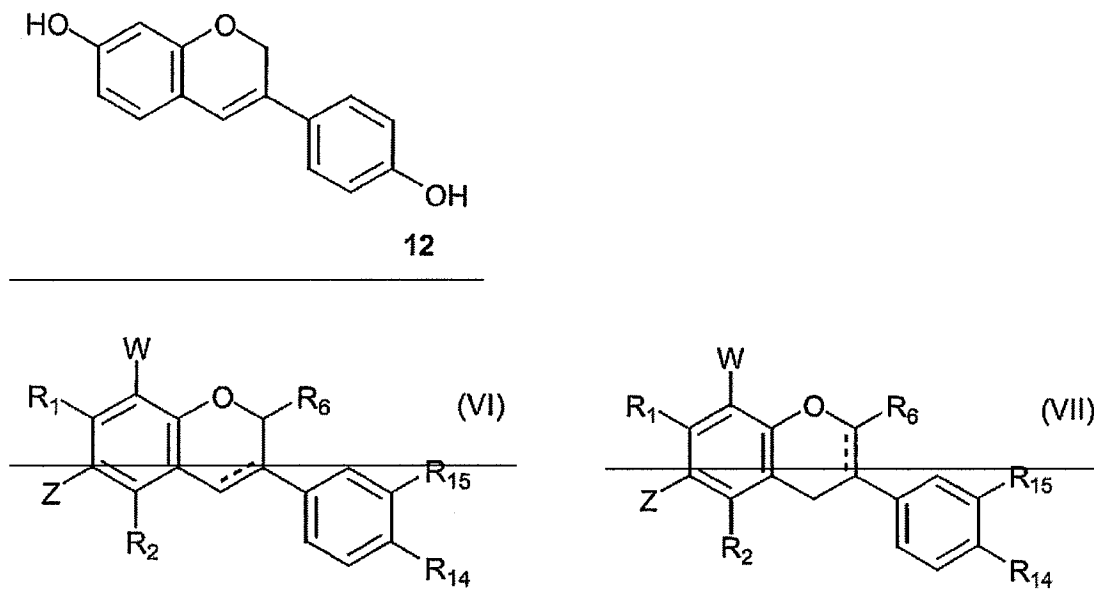


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

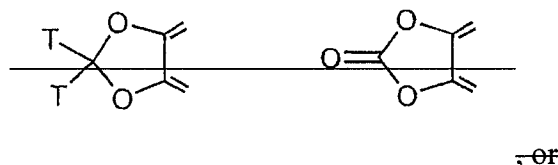
LISTING OF CLAIMS:

1. (currently amended): A method of increasing the sensitivity of cancer cells or a tumour to a chemotherapeutic agent by contacting said cells or tumour with an isoflavonoid compound of formula 12(VI) or (VII):

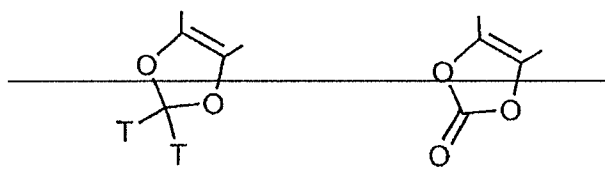


wherein

~~R₁, R₂ and Z are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₂ is as previously defined, and R₁ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from~~



~~R₁ is as previously defined, and R₂ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from~~



~~W is R₁;~~

~~R₃ is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R₁₁ where R₁₁ is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO₂R₁₂ where R₁₂ is hydrogen, alkyl, haloalkyl, aryl or arylalkyl;~~

~~R₄ is hydrogen, alkyl or aryl, or~~

~~R₃ and R₄ taken together with the nitrogen to which they are attached comprise pyrrolidiny or piperidiny;~~

~~R₆ is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR₃R₄, COR₁₁ where R₁₁ is as previously defined, CO₂R₁₂ where R₁₂ is as previously defined or CONR₃R₄;~~

~~R₉ is alkyl, haloalkyl, aryl, arylalkyl, C(O)R₁₁ where R₁₁ is as previously defined, or Si(R₁₃)₃ where R₁₃ where each R₁₃ is independently hydrogen, alkyl or aryl;~~

~~R₁₀ is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino;~~

~~the drawing “—” represents either a single bond or a double bond;~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R₁₄ and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀,
COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl,
thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused
together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

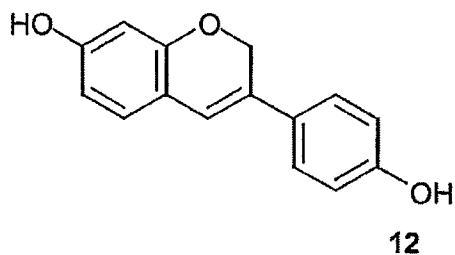
~~wherein the cancer is ovarian, pancreatic or prostate cancer, and~~

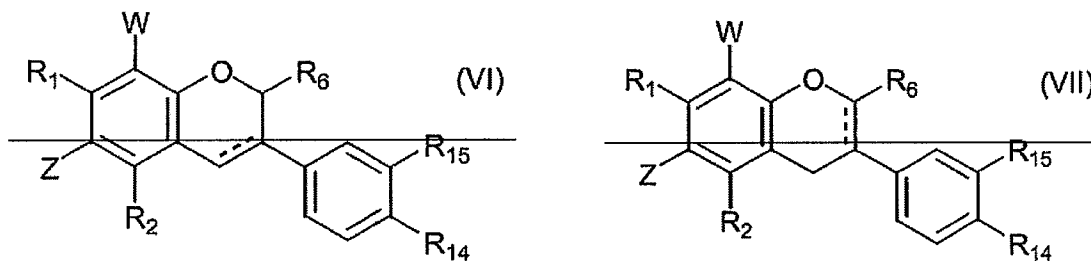
~~the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin,
paclitaxel, gemcitabine or doxorubicin.~~

2. (previously presented): A method of claim 1, wherein prior to the contacting, the cancer cells or tumour were/was not sensitive to the chemotherapeutic agent.

3. (currently amended): A method of claim 1, wherein the compound of formula 12(VI) or (VII) is administered to a subject in need of such treatment.

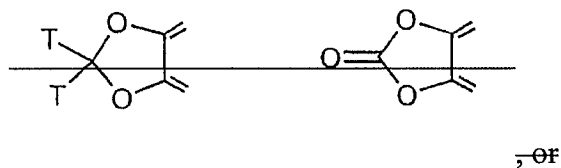
4. (currently amended): A combination therapy for the treatment or prophylaxis of ~~cell proliferation, cancer or a disease associated with oxidant stress~~ comprising administering to a subject a therapeutically effective amount of a compound of formula 12(VI) or (VII) and a chemotherapeutic agent:



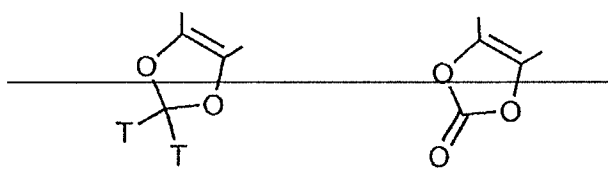


wherein

~~R₁, R₂ and Z are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₂ is as previously defined, and R₁ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from~~



~~R₁ is as previously defined, and R₂ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from~~



W is R₁;

~~R₃ is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R₁₁ where R₁₁ is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO₂R₁₂ where R₁₂ is hydrogen, alkyl, haloalkyl, aryl or arylalkyl,~~

~~R₄ is hydrogen, alkyl or aryl, or~~

~~R₃ and R₄ taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl,~~

~~R₆ is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR₃R₄, COR₁₁ where R₁₁ is as previously defined, CO₂R₁₂ where R₁₂ is as previously defined or CONR₃R₄,~~

~~R₉ is alkyl, haloalkyl, aryl, arylalkyl, C(O)R₁₁ where R₁₁ is as previously defined, or Si(R₁₃)₃ where R₁₃ where each R₁₃ is independently hydrogen, alkyl or aryl,~~

~~R₁₀ is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino,~~

~~the drawing “—” represents either a single bond or a double bond,~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R₁₄ and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

~~wherein the cancer is ovarian, pancreatic or prostate cancer, and~~

~~the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicin.~~

5.-7. (canceled).

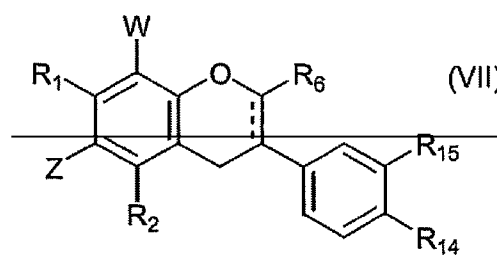
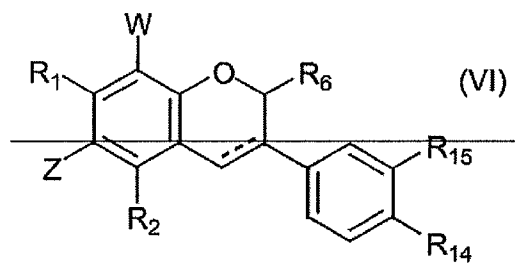
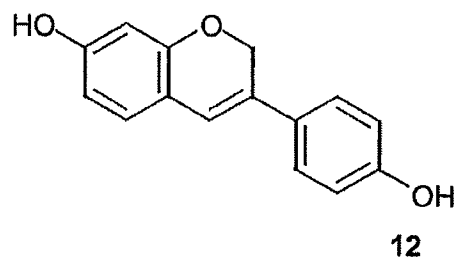
8. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) ~~or~~ (VII) precedes the administration of the chemotherapeutic agent.

9. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) ~~or~~ (VII) and the chemotherapeutic agent is simultaneous.

10. (currently amended): A method claim 4, wherein the combination therapy follows observed resistance by cancer cells or tumour to ~~[[a]]~~the chemotherapeutic agent.

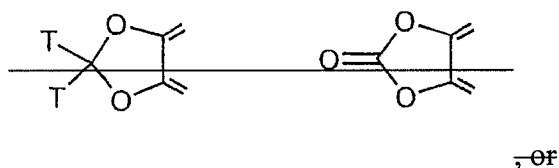
11.-22. (canceled).

23. (currently amended): A pharmaceutical composition comprising a compound of formula 12(VI) ~~or~~ (VII) and a chemotherapeutic agent:

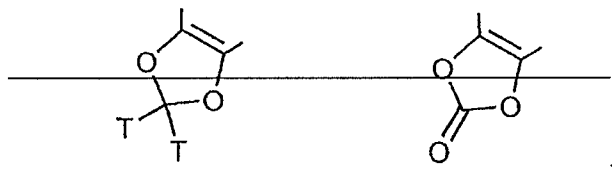


wherein

~~R₁, R₂ and Z are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀,
 COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl,
 alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or
 R₂ is as previously defined, and R₁ and Z taken together with the carbon atoms to which they are
 attached form a five-membered ring selected from~~



~~R₁ is as previously defined, and R₂ and Z taken together with the carbon atoms to which they are
 attached form a five-membered ring selected from~~



~~W is R₁;~~

~~R₃ is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R₁₁ where R₁₁ is hydrogen,
 alkyl, aryl, arylalkyl or an amino acid, or CO₂R₁₂ where R₁₂ is hydrogen, alkyl, haloalkyl,
 aryl or arylalkyl;~~

~~R₄ is hydrogen, alkyl or aryl, or~~

~~R₃ and R₄ taken together with the nitrogen to which they are attached comprise pyrrolidinyl or
 piperidinyl;~~

~~R₆ is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR₃R₄, COR₁₁ where R₁₁ is as previously
 defined, CO₂R₁₂ where R₁₂ is as previously defined or CONR₃R₄;~~

~~R₉ is alkyl, haloalkyl, aryl, arylalkyl, C(O)R₁₁ where R₁₁ is as previously defined, or Si(R₁₃)₃~~

~~where R₁₃ where each R₁₃ is independently hydrogen, alkyl or aryl,~~

~~R₁₀ is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino,~~

~~the drawing “—” represents either a single bond or a double bond,~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R₁₄, and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀,~~

~~COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl,~~

~~thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused~~

~~together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

~~wherein the chemotherapeutic agent is platinum-based or anti-mitotic agent~~cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicins.

24.-28. (Canceled).